MANAGED PROBLEM SOLVING (MAPS)

Good Evidence – Medication Adherence

INTERVENTION DESCRIPTION

Target Population

• HIV clinic patients who are antiretroviral treatment experienced or naïve

Goals of Intervention

Improve adherence to antiretroviral therapy

Brief Description

MAPS is an individual-level, problem-solving intervention delivered in person and via telephone calls to HIV clinic patients. The intervention focuses on improving medication adherence through an iterative, five-step process which consists of 1) identifying barriers to adherence, 2) brainstorming to generate potential solutions, 3) decision-making and developing a plan of action, 4) implementing the plan, and 5) evaluating and modifying the plan as necessary. In-person sessions include education related to the treatment regimen and to common medication misperceptions; problem-solving to identify daily routines, cues, cognitive aids and social supports; screening to identify barriers related to depression, substance use, toxicity management and competing demands; and review of adherence data to determine where problems have occurred and to develop solutions. In addition, on-going telephone calls reinforce content delivered during the in-person sessions, allow for additional problem solving, remind patients to obtain refills, and encourage continued adherence to intervention strategies.

Theoretical Basis

Social Cognitive Theory

Intervention Duration

• Four in-person sessions: one 60-90 minute session followed by three 20-45 minute sessions delivered monthly over 3 months; twelve weekly telephone calls (20-30 minutes per call) during the first 3 months, and monthly telephone calls throughout the remaining 9-month intervention period

Intervention Setting

HIV clinics

Deliverer

• Trained interventionists, with at least a college degree and some prior experience working with a patient population

Delivery Methods

- Discussion
- Goal setting/plan

- Lecture/teach
- Memory aids/cues

Problem solving

INTERVENTION PACKAGE INFORMATION

The intervention manual is available for download at

http://www.med.upenn.edu/cceb/maps-form.shtml

EVALUATION STUDY AND RESULTS

The original evaluation was conducted in Philadelphia, Pennsylvania between 2005 and 2010.

Key Intervention Effects

- Increased medication adherence
- · Achieved undetectable viral load

Study Sample

The baseline study sample of 180 men and women living with HIV is characterized by the following:

- 85% black or African American, 13% white, 2% other
- 60% male, 40% female
- Mean age of 43 years, range: 19-65 years
- 40% treatment-naïve, 60% treatment experienced

Recruitment Settings

HIV clinics

Eligibility Criteria

HIV clinic patients were eligible if they were at least 18 years of age, had an HIV-1 plasma viral load >1000 copies/mL, and were either 1) treatment naïve and initiating a standard regimen or 2) treatment experienced and either restarting their most recent suppressive regimen or initiating a new regimen.

Assignment Method

HIV clinic patients (N = 180) were randomly assigned to 1 of 2 study arms: MAPS (n = 91) or usual care comparison (n = 89).

Comparison Group

The usual care comparison included meetings with a pharmacist for regimen education and, if desired, provision of pill organizers.

Relevant Outcomes Measured and Follow-up Time

- Medication adherence behavior (recorded using MEMs caps) was defined as the proportion of prescribed doses taken. Medication adherence was categorized as ≤70%, 71-80%, 81%-90%, 91-95%, or >95% and was assessed at 3, 6, 9, and 12 months post-initiation of intervention.
- Viral load was measured at 3, 6, 9 and 12 months post-initiation of intervention and was assessed as undetectable (<75 copies/mL) or detectable.

Participant Retention*

- MAPS Intervention
 - o 75% retained at 3 months post-initiation of intervention
 - o 68% retained at 6 months post-initiation of intervention
 - o 76% retained at 9 months post-initiation of intervention
 - o 84% retained at 12 months post-initiation of intervention
- Usual Care Control
 - o 75% retained at 3 months post-initiation of intervention
 - o 79% retained at 6 months post-initiation of intervention
 - o 78% retained at 9 months post-initiation of intervention
 - o 87% retained at 12 months post-initiation of intervention

Significant Findings

- Across the four assessments (3 to 12 months post-initiation of intervention), intervention participants were significantly more likely to be in a higher adherence category than comparison participants (OR = 1.78, 95% CI = 1.07 2.96, missing data imputed; Adj OR = 2.33, 95% CI = 1.35 4.05, without imputation).
- Across the four assessments (3 to 12 months post-initiation of intervention), intervention participants were significantly more likely to have an undetectable viral load (<75 copies/mL) than comparison participants (Adj OR = 1.98, 95% CI = 1.15 -3.41, without imputation).

Considerations

- This study did not meet the best-evidence criteria due to <70% retention rate per arm at each assessment and no significant intervention effects on the undetectable viral load outcome across the four assessments in an intention-to-treat analysis with missing data imputed.
- At baseline, 69% were on a protease inhibitor (PI)-based regimen and 28% were on a nonnucleoside reverse transcriptase inhibitor (NNRTI)-based regimen.
- The MEMs cap was placed on one medication bottle; for patients receiving multiple antiretroviral drugs, the monitored drug was selected in the following order of preference: 1) Nonnucleoside analog reverse transcriptase inhibitor, 2) protease inhibitor (ritonavir first), 3) integrase inhibitor, 4) entry inhibitor, or 5) nucleoside reverse transcriptase inhibitors.

REFERENCES AND CONTACT INFORMATION

Gross, R., Bellamy, S. L., Chapman, J., Han, X., O'Duor, J., Palmer, S. C., . . . Strom, B. L. (2013). <u>Managed problem solving for antiretroviral therapy adherence: A randomized trial</u>. *JAMA Internal Medicine*, *173*, 300-306.

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